Signal transductio Full-length rat an

Signal transductio Human androgen rec DNA encoding novel Andorgen receptor Fused androgen rec Human androgen rec Human androgen rec Human immune syste

Human DNA for stag

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Green fluorescent Colon adenocarcino

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ALIGNMENTS

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Human; AIB1; amplified in breast cancer 1; androgen receptor; AR; prostate cancer; chromosome X;\ ds.
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/product= "Human androgen receptor (AR) protein"
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1115..3874
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Human androgen receptor (AR) gene.
                                                                                                                                                                                                                                                                                                                                                   AAD30440 standard; DNA; 4321 BP.
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Query Match

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The invention relates to a method for assessing the risk of prostate cancer in a human subject. The method involves determining the length of the contiguous CAG or CAA repeats in both AIDI (Amplified in Breast cancer 1) gene alleles or contiguous CAG, CAA or GGN repeats in the androgen receptor gene of the subject. The method is useful for assessing a subject s risk for acquiring or developing prostate cancer. The present sequence is human androgen receptor (AR) gene. Human AR gene is located on X chromosome. Assessing the risk of acquiring or developing prostate cancer in a human subject, comprises determining the length of the contiguous CAC, CAA and/or GGN repeats in the AIBI gene and/or androgen receptor gene of the subject -Disclosure; Page 61-62; 86pp; English

4321 BP; 966 A; 1281 C; 1168 G; 906 T; 0 other;

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AGCTGCTAAAGACTCGGAGGAAGCAAGGAAAGTGCCTGGTAGGACTGACGGCTGCCTTTG 1046 1166 1226 1285 1345 1405 1465 358 418 478 537 597 657 986 238 298 813 873 931 Gaps 59 CTTTGGCTGCGAGCGGGCGAGNCTAGCTGCACATTGCAAAGAAGGCTCTTAGGAG-CAGG CGACTGGGGAGCGGCTTCAGCACTGCAGCCACGACCNGCCTGGTTAGGCTGCACGCGGAG CGAA-GGACGCACCACCCAGCCCCAGCTCCAGCGACAGCNAACGCCTCTTGCA-CGAGATCCCGGGGAGCCAGCTTGCTGGGAGAGCGGGAACGGTCCGGAGCAAGCCCAGAGG CAGAGGAGGCGACAGAGGGCAAAAAGGGCCCNAGCTAGCCGCTCCAGTGCTGTACAGNAGC DB 24; Length 4321; 24; Indels 16; Score 4188.2; Pred. No. 0; 0; Mismatches Query Match 82.4%; Best Local Similarity 99.1%; Matches 4297; Conservative 1346 1286 1406 479 538 598 419 658 754 09 874 932 179 239 1047 299 1107 359 1167 1227 987 g ò a ŏ g δy g q οy g 9 2 2 QΥ q ΟY g οy 셤 δλ ογ ò g ð

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AGGC AGGC	CTTCCAGCACCACCACCCACTGAGGAGACAAACCCAGAAGCTGACAGTGTCACACA 3865
TTG2 TTG2	TTTCTGAATGTCCTGGAAGCCATT3AGCCAGGTG 392.
$\omega - \omega$	IGTGTGGTGGACACGACAACAACCAGCCCGACTCCTTTGCAGCCTTGCTGTTTAGCC 398
TCA TCA	GTACACGTGGTCAAGTGGGCCAAGGCCTTGCCTG 404
GCT	TCCGCAACTTACACGTGGACGACCAGATGGCTGTCATTCAGTACTCCTGGATGGGGC 410
TCATO TCATO	3GTGTTTGCCATGGGCTGGCGATCCTTCACCAATGTCAACTCCAGGATGCTCTACT 416
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GTG GTG	TCCGAATGAGGCACCTCTCTCAAGAGTTTGGATGGCTCCAAATCACCCCCAGGAAT 428
TCCT	3TGCATGAAAGCACTGCTACTCTTCAGCATTATTCCAGTGGATGGGCTGAAAAATC 43.
AAAAA AAAAA	TCTTTGATGAACTTCGAATGAACTACATCAAGGAACTCGATCGTATCATTGCAT 440
GCA.	AAAAAATCCCACATCCTGCTCAAGACGCTTCTACCAGGTCACCAAGCTCCTGG 446
ACT ACT	CCGTGCACCTATTGCGAGAGCTGCATCACTTCTGACCTGCTAATCAAGT 4525
CAC	CCATGGTGAGCGTGGACTTTCCGGAAATGATGGCAGAGATCATCTCTGTG:AAGTGC 45E
CAA CAA	GATCCTTTCTGGGAAAGTCAAGCCATCTATTCCACACCCAGTGAAGGTTGGAA 46.
5 = 5	CTATTTCCCACCCCAGGTCATGCCCCTTTCAGATGTCTTCTGCCTGTTATAACTC 4705
GCA GCA	CTACTCCTCTGCAGTGCCTTGGGGAATTTCCTCTATTGATGTACAGTCTCTCATGA 476

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Proapoptotic; dependence domain; p75NTR; androgen receptor; DCC; huntingtin polypeptide; Machado-Joseph disease; SCA1; SCA2; SCA6; atrophin-1; cell death; apoptosis; Huntington's disease; head trauma; Alzheimer's disease; Kennedy's disease; spinocerebellar ataxia; stroke; dentatorubropallidoluysian atrophy; cell proliferation; cell survival; neoplastic; malignant; autoimmune; fibrotic; ss.
                                                                                                                        This invention describes novel pure proapoptotic dependence peptides which comprise a sequence of an active dependence domain selected fro dependence polypeptides consisting of p75NTR, androgen receptor, DCC,
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TCTTCCCTCCCTATCTAACCCTCCCATGGCACCTTCAGACTTTGCTTCCCATTGTGGCTC
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huntingtin polypeptide, Machado-Joseph disease gene product, SCA1, SCA2, SCA6 and atrophin-1 polypeptide. The proapoptotic peptides are capable of inducing cell death and can be used to develop products to mediate or inhibit apoptosis. The methods can be used for reducing the severity of a proapoptotic dependence domain mediated pathological conditions e.g. Huntington's disease, Alzheimer's disease, Kennedy's disease, Spinocerebellar ataxias, dentatorubropallidoluysian atrophy. Machado-Joseph disease, stroke or head trauma. They can also be used for reducing the severity of a pathological condition mediated by upregulated eell proliferation or cell survival e.g. neophastic, malignant, autoimmune or fibrotic conditions. This sequence encodes a human androgen receptor described in the method of the invention.
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Diagnosing and detecting the progression of liver cancer, hepatocellular carcinoma or metastatic liver tumor in a patient, involves detecting the level of expression of two or more genes

Diagnosing and detecting hepatocellular carcinoma

liver tissue sample

Vockley JG;

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Peres-Da-Silva

Horne D, Alvares C,

WPI; 2002-426119/45.

02-OCT-2000; 2000US-237054P. 02-OCT-2001; 2001WO-US30589

LOGIC INC

(GENE-) GENE

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Gene; liver cancer; ds; hepatocellular carcinoma; hepatotropic; metastatic liver tumour; cytostatic; expression profile; disease state; disease progression; drug toxicity; drug efficacy; drug metabolism.
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                                ACTCGATCGTATCATTGCATGCAAAAAAAATCCCACATCCTGCTCAAGACGCTTCTA
                                       Gene #2300 used to diagnose liver cancer.
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The invention relates to a novel method for diagnosing and detecting the progression of liver cancer, hepatocellular carcinoma or metastatic liver cancer from tumour in a patient, and differentiating metastatic liver cancer from chaptocellular carcinoma in a patient, involving detecting the level of expression of two or more genes represented in ABN3503-ABN37455 in a expression of two or more genes represented in ABN3503-ABN37455 in a carcinoma sample. The method is useful for diagnosing and detecting cytostatic activity. The method is useful for diagnosing and detecting the progression of liver cancer, hepatocellular carcinoma and metastatic liver carcinoma in a patient. The method is useful for identifying expension profiles which serve as useful diagnostic markers as well as markers that can be used to monitor disease states, disease progression, drug toxicity, drug efficacy and drug metabolism.

Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO cat fit, wipo.int/pub/published_pct_sequences.
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Rest Local Similarity 98.6%; Pred. No. 0;
Matches 3685; Conservative 0; Mismatches
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3074 3134 1778 CGGTICIGGGICACCCICAGCCGCCGCTICCTCATCCTGGCACACTCTCTTCACAGCCGA 3194 3245 3305 3014 1898 1958 3425 2078 3485 2138 3545 2198 3605 2258 3665 2318 3725 2378 3785 2438 3845 2498 3962 2618 4025 2678 CGACTACTACAACTITCCACTGGCTCTGGCCGGACCGCCGCCCCCTCCGCCGCCTCCCCA AGAAGGCCAGTTGTATGGACCGTGTGGTGGTGGGGGGTGGT------GGCGGCGG 3366 CGCACCTGATGTGTGTACCCTGGCGGCATGGTGAGCAGAGTGCCCTATCCCAGTCCCAC 2079 TICTGTCAAAAGCGAAATGGGCCCCTGGATGGATAGCTACTCCGGACCTTACGGGGACAT TTGTGTCAAAAGCGAAATGGGCCCCTGGATGGATAGCTACTCCGGACCTTACGGGGACAT GACCTGCCTGATCTGTGGAGATGAAGCTTCTGGGTGTCACTATGGAGCTCTCACATGTGG AAGCTGCAAGGTCTTCTTCAAAAGAGCCGCTGAAGGGAAACAGAAGTACCTGTGCGCAG GAAGCTGACAGTGTCACACATTGAAGGCTATGAATGTCCAGCCCCATCTTTC1GAATGTCCT GGAAGCCATTGAGCCAGGTGTAGTGTGTGCTGGACACGACAACAACAACCAGCCCGACTCCTT 1599 2955 3015 1659 3075 1719 3135 1779 3195 1839 3426 3486 2139 3546 2199 2259 2319 3726 3606 3666 2379 3786 2439 3846 2499 3906 2559 2619 3966 qq qq δý δ QQ Q മ : Ω Qy Db QY Db δŻ q Οy g δy δy g Qγ Ω οy Ωp Qγ qq δÿ οg Qγ q δ g Qγ qq Ωy a

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	2715 TCCTTG	1359 TCCTTG	2775 CACTGA 1419 CACTGA	ro or	2895 GTCTAC 1539 GTCTAC	2955 CGACTA 1599 CGACTA	3015 TCCCCA 1659 TCCCCA	3075 GGCGGC 1719 GGCGGC	3135 CGGTTC7 1779 CGGTTC7	3195 AGAAGGC 1839 AGAAGGC	3246 CGGCGGC 1899 CGGCGGC	3306 AGCCCC 	3366 CGCACCT 2019 CGCACCT	3426 TTGTGTC 2079 TTGTGTC	3486 GCGTTTG 2139 GCGTTTG	3546 GACCTGC 2199 GACCTGC	3606 AAGCTGC 2259 AAGCTGC	3666 CAGAAAT 2319 CAGAAAT	3726 GAAATGT 2379 GAAATGTT
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Db 243 TTTTAAGATCTGGGCATCTTTGAATCTACCCTTCAAGTATTAAGAGACAGAGTGTGAGG	STCA 16	Db 303 CTAGCAGGCAGATCTTGTCCACGTGTGTCTTCTGCACGAGACTTTGAGCCTGTCA 362	1695 GAGCGCTTTTTGCGGGGTTGCTCCCGCAAGTTTCCTTCTTGGACCTTCCCGCAGGTGGG 1	1755 CAGCTAGCTGCACTACCGCATCATCATCATTACTTTCTTCTCTGCAGGTGGG 4	1815	1875 GTTAGGGCTGGGAAGGTCTACCCTCGGCGGCGCGCGAGACCTACCGAGGAGCTTCCA	1935 GAATCTGTTCCAGAGCGTGCGCAAAGTGATCCAGAACCCGGGCCCCAGGCACCCAGAGGC	1995 CGCGAGCGCAGCTCCCGGCGCCAGTTTGCTGCTGCTGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAG	2055 GCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAG	2115 TAGCCCCAGGCAGCAGCAGCAGCAGGTGAGGATGTTCTCCCCAAGCCCATGG 2	2175 AGGCCCACAGGCTACCTGGTCCTGGATGAGGAACAGCAACCTTCACAGCCGCAGTCGC 22 111111111111111111111111111111111	2235 CCTGGAGTGCCACCCGAGAGGTTGCGTCCCAGAGCCTGGAGCCGCCGTGGCCGCCA 2 1111111111111111111111111111111111	23	24 24 10	24	253	259	265	271
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GCAAGAG 27 GCAAGAG 14	CACCAAAGGCTAGAAGG 2834	3GACACTTGAACTGCC 28 	TGCGTACCAGAGTCG 29	36CCGCCTCCCCA 3	CTGGGGGGTGC 307.	SGGACC 3	CTTCACAGCCGA 319. 	GGCGG 324	AGCTGT 330:	TTCAC 336.	CCCAC 342	GACAT 348	TTCCACCCAGAA 354 	CTCTCACATGTGG	AGTACCTGTGCGCCA	CTTGTCGTCTTCG 3	BAAACTTGGTAATCT 3
CCTTGTGCCCCATTGGCCGAATGCAAAGGTTCTCTGCTAAA 	CTGAAGATACTGCTGAGTATTCCCCTTTCAAGGAGGTA 	AGAGCCTAGGCTGCTCGGCAGCGCTGCAGCAGGAGGAGCT 	CTACCCTGTCTCTACAAGTCCGGAGCACTGGACGAGG 	ACTACTACAACTTTCCACTGGCTCTGGCCGGACCGC 	CCCACGCTCGCATCAAGCTGGAGAACCCGCTGGACT	GGGGGAGTGCGGTAFGGGAACCTGGCGAGCTGG 	STICTGGGTCACCCTCAGCGGCGCTTCCTCATCCT 	GCCAGTTGTATGGACCGTGTGGTGGTGGTGGTGGTGTT	39CGGCGGCGGCGGCGGCGGCGCGCGCGCGCGCGCGCGCG	GGCCCCTCAGGGGCTGGC 	ATCTGTGGTACCCTGCGGCATGGTGAGCAC 	aaagcgaaatgggccctggatggatagctb 	ITTGGAGACTGCCAGGGACCATGTTTTGCCCATTGACTAI 	CIGCCIGATCIGIGGAGAIGAAGCITCIGGGIGTCACTAI 	CTGCAAGGTCTTCTTCAAAAGGCCGCTGAAGGGAAACAC 	AATGATTGCACTATTGATAAATTCCGAAGGAAAAATTGT 	rgttatgaagcagggatgactctgggaggggaagctg
2715 TC 1359 TC	2775 CA 1419 CA	2835 CG 1479 CG	2895 GT 1539 GT	2955 CG 1599 CG	3015 TC	3075 GG(1719 GG(3135 CG(1779 CG(3195 AG 1839 AG	3246 CG(1899 CG(3306 AGC 	3366 CGC 2019 CGC	3426 TTG 2079 TTG		3546 GAC	3606 AAG 2259 AAG	3666 CAG 2319 CAG	1726 GAAA
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11111111111111111111111111111111111111	4926 TGATGATCCTCATATGGCCCAGTGTCAAGTTGTGCTTGTTACAGCACTACTCTGTGCCA 4985	4986 GCCACACAAACGTTTACTTATGTTATGCCACGGGAAGTTTAGAGAGCTA [†] 	5046		RESULT 5 AAT63407 ID AAT63407 standard; cDNA; 3569 BP. , , , , , , , , , , , , , , , , , , ,	AAT63407; 22-JUN-1997 (first entry)	Androgen receptor cDNA. Androgen receptor; acidic fibroblast growth factor; aFGE; Androgen receptor; acidic fibroblast growth factor; therapy; antisense; benign prostatic hyperplasia; prostate cancer; therapy;		Key CDS misc_feature	/*tag= /note= " misc_feature compleme	misc_feature		WO9711170-A1. 27-MAR-1997.	20-SEP-1996; 96WO-US15081. 20-SEP-1995; 95US-0004018.			PT Oligonucleotide(s) antisense to numan amurogen receptor mineral properties of properties of prostatic hyperplasia PT FGF genes - used to inhibit gene expression, for the treatment of properties		oligonucleotides (see also AAT63200, AAT63404-03) to tils
Db	QY Db	QQ Dp	οy	qa	2212	2×2;	X X X E	Z X O X	(E. E. E. E.	चित्र चित्र : :	E E E E	H H H X	, , , , , ,						
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3786 GAAACTACAGGAGGAGGAGAGGCTTCCAGCACCACCAGCCCCACTGAGGAGACAACCCA 3845	2439 GAAACTACAGGGGGGGGGGGGGGGCTCCAGCACCACCAGCCACCAGGAGACAACCCA 2498 3846 GAAGCTGACAGTGTCACACACATAGAGGCTATGAATGTCAGCCCATCTTCTGAATGTCT 3905	2499 GAAGCTGACAGTGTCACACATTGAAGGCTATGAATGTCACCCCATCTTTCTGAATGTCCT 2558 3906 GGAAGCCATTGAGCCAGGTGTGTGTGTGTGTGTGACACGACAACAACCAGCCCGACTCTT 3965	2559 GGAAGCCATTGAGCCAGGGTGTGTGTGTGTGGACCAGACCAGCCCGACTCCTT	3966 IGCAGCCTTGCTCTCTAGCCTCAATGAACTGGGAGAGAGACAGCTTGTACACGTGGTCAA 4025 111111111111111111111111111111111111	4026	4086	4146	4206 GICCCGGAIGTACAGCCAGIGIGCCGAAIGAGGCACCICTCAAGAGIITGGAIGGCT 4265 111111111111111111111111111111111111	4266	4326	4386 ACTCGATCGTATCATTGCATGCAAAAAAAATCCCACATCCTGCTCAAGACGCTTCTA 4445	Y 4446 CCAGCTCACCAAGCTCCTGGACTCCGTGCAGCCTATTGCGAGAGAGCTGCATCAGTTCAC 4505	4506	4566	OY 4626 CACCCAGIGAAGCATIGGAAACCCTATITCCCCACCCCAGTCAIGCCCCCTITCAGAIG 4685 DD 3279 CACCCAGIGAAGCAIIGGAAACCCTATITCCCCACCCAGCICAIGCCCCCTITCAGAIG 3338	4686	TTTTCTC 	QY 4806 TTTCTCTCTTTTTTTTTTTTTCTTCCTCCTATCTAACCCTCCCATGGCACCTTCAGAC 4865	QY 4866 TITGCTICCCAIIGIGGCICCTAICIGIGITITGAAIGGIGITGIAIGCCTITAAAICIG 4925
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receptor sequence or antisense oligonucleotides (see also AAT63406) to the human acidic fibroblast growth factor gene (see also AAT63197-99). The methods are esp. useful for the treatment of benign prostatic hyperplasia and prostate cancer.
                                                                                                                                    TCCCGCAGGTGGGCAGCTAGCTGCAGCGACTACCGCATCATCACAGCCTGTTGAACTCTT
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2581 2761 2701 1188 2941 1428 3121 3241 3181 1728 948 1908 3481 3541 2028 CCGCTGACCTTAAAGACATCCTGAGCGAGGCCAGCACCATGCAACTCCTTCAGCAACAGC CCACTTCCTCCAAGGACAATTACTTAGGGGGCACTTCGACCATTTCTGACAACGCCAAGG AGTTGTGTAAGGCAGTGTCGGTGTCCATGGGCCTGGGTGTGGAGGCGTTGGAGCA.?CTGA GTCCAGGGGAACAGCTTCGGGGGATTGCATGTACGCCCCACTTTTGGGAGTTCCACCCG CTGTGCGTCCCACTCCTTGTGCCCCATTGGCCGAATGCAAAGGTTCTCTGCTAGACGACA GCGCAGGCAAGAGCACTGAAGATACTGCTGAGTATTCCCCTTTCAAGGGAGGTfACACCA AAGGGCTAGAAGGCGAGAGCCTAGGTGCTGGCAGCGCTGCAGCAGGAGGTCCGGGA CACTTGAACTGCCGTCTACCTGTCTCTACAAGTCCGGAGCACTGGACGAGGCAGCTG CGTACCAGAGTCGCGACTACTACAACTTTCCACTGGCTCTGGCCGGACCGCCCCCCTC CGCCCCCCCCATCCCCACCGCTCGCATCAAGCTGGAGAACCCGCTGGACTACGGCAGCG CCTGGGCGGCTGCGGCGCGCAGTGCCGCTATGGGGAACCTGGCGAGCCTGCATGGCGCGG GTGCAGCGGGACCCGGTTCTGGGTCACCCTCAGCCGCCGCTTCCTCATCCTGGCACACTC TCACCGCACCTGATGTGTGTACCCTGGCGGCATGGTGAGCAGGTGCCCTATCCCAGTC CCACTTGTGTCAAAAGCGAAATGGGCCCCTGGATGGATAGCTACTCCGGACCTTACGGGG ACATGCGTTTGGAGACTGCCAGGGACCATGTTTTGCCCATTGACTATTACTTTCCACCC 889 2462 949 2522 1009 2582 1069 2642 2702 2762 2822 2882 1369 2942 1489 3062 1549 1429 3002 3122 1609 3182 1669 3242 1729 3302 3362 1849 1789 3422 1909 3482 1969 δ qq g οy ò QQ Ω Db ò ρp δy Q ò qq οy Db Ω Ωp δy g ò QQ qq Ω QΥ g Qγ g òγ g ÓΥ g δ g ŏ g

	4 4 3 2 1	2 GECTCCAAATCACCCCCAGGAATTCCTGTGCATGAAAGCACTGCTACTTTCAGCATTT 2 GECTCCAAATCACCCCCCAGGAATTCCTGTGCATGAAAGCACTGCTACTCTTCAGCATTT 9 GECTCCAAATCACCCCCCAGGAATTCCTGTGCATGAAAGCACTGCTACTCTTCAGCATTTTTTTT	2 7 8 8 8 8 4 9 7 5 5
	280 438 286	9 GGCTCCAAATCACCCCCCAGGAATTCCTGTGCATGAAAGCACTGCTACTTCAGCAT 2 TTCCAGGATGGGATGGAAAAATTCATATGATGAAGAACTTCGAATGAACTACAT 2 TTCCAGTGGATGGGTGGAAAAATTCAAAAATTCTTTGATGAACTTCGAATGAACTACAAT 3 TTCCAGTGGATGGGCTGAAAAATCAAAAATTCTTTGATGAACTTCGAATGAAT	N 8
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	4261 2748	2 ACAAGTCCCGGATGTACAGCCAGTGTGTCCGAATGAGGCACCTCTCTCAAGAGTTTGGAT 	420
	420	2 AIGTCAACTCCAGGATGCTCTACTTCGCCCCTGATCTGGTTTTCAATGAGTACCGCATGC	414
٠	4141 2628	2 TCATTCAGTACTCCTGGATGGGGCTCATGGTGTTTGCCATGGGCTGGCGATCCTTCACCA	408
•	4081	2 TCAAGTGGGCCAAGGCCTTGCCTGGCTTCCGCAACTTACACGTGGACGACCAGATGGCTG	402
	4021	2 CCTTTGCAGCCTTGCTCTCTAGCCTCAATGAACTGGGAGAGAGA	396:
	3961 2448	2 TCCTGGAAGCCATTGAGCCAGTGTAGTGTGTGCTGGACACACAACAACAACCAGCCCGACT	3903
	3901	2 CCCAGAAGCTGACAGTGTCACACTTGAAGGCTATGAATGTCAGCCCATCTTTCTGAATG	3842
	3841	ATCTGAAACTACAGGAGGAAGGAGGCTTCCAGCACCACCAGCACCCCCTGAGGAGACAA 	3782
	3781 2268	TTCGGAAATGTTATGAAGCAGGGATGACTCTGGGAGCCCGGAAGCTGAGAAAAACTTGGTA 	3722
	3721 2208	CCAGCAGAAATGATTGCACTATTGATAAATTCCGAAGGAAAAATTGTCCATCTTGTCGTC 	3662
٠.	3661 2148	GIGGAAGCIGCAAGGICITCITCAAAAGAGCGGAAGGGAAACAGAAGTACCIGGGG 	3602
	3601 2088	AGAAGACCTGCCTGATCTGTGGAGATGAAGCTTCTGGGTGTCACTATGGAGCTCTCACAT	3542

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Complementary DNA sequences derived from the cDNA may be used as probes
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                                                                                                                                                        4982 GCCAGCCACACAAACGTTTACTTATCTTATGCCACGGGAAGTTTAGAGAGGTAAGATTAT
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            DNA encoding androgen receptor protein - useful for transforming eukaryotic hosts for protein expression and subsequent antibody protein expression and subsequent
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                                                                                                                                                                                                                                                                        AAN91772 standard; cDNA; 3569
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P-PSDB; AAP93109.
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Fri May

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C de Good See C	tect the presence of androgen rece tect AR gene defects using DNA hyb	uence 3509 BP; 796 A; 1008 C; 975 Match 69.8%; Score 3 Cocal Similarity 99.6%; Pred. M	7 7	2 TTTTGTTTT 	2 ACAGACTGT 	2 TTTGAG 	2 TCCCGC 1 TCCCGC			7 7		7 7	C) Ch	0 0				
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2581 2641 1068 GTCCAGGGGAACAGCTTCGGGGGATTGCATGTACGCCCCACTTTTGGGAGTTCCACCG 2701 2881 2941 2521 1428 3001 1488 3061 1548 3121 1608 3181 1668 3241 1728 3301 3361 1848 1908 3421 3481 3541 2028 3542 AGAAGACCIGCCIGATCIGIGGAGAIGAAGCIICIGGGIGICACIAIGGAGCICICACAI 3601 **AGTTGTGTAAGGCAGTGTCGGTGTCCATGGGCCTGGGTGTGGAGGCGTTGGAGCATCTGA** CCACTTCCTCCAAGGACAATTACTTAGGGGGCACTTCGACCATTTCTGACAAGGCCAAGG GCGCAGGCAAGACACTGAAGATACTGCTGAGTATTCCCCTTTCAAGGGAGGTTACACCA CACTTGAACTGCCGTCTACCCTGTCTCTACAAGTCCGGAGCACTGGACGAGGCAGCTG 1369 CACTTGAACTGCCGTCTACCCTGTCTCTACAAGTCCGGAGCACTGGACGAGGCAGCTG AAGGGCTAGAAGGCGAGAGCCTAGGCTGCTCTGGCAGCGCTGCAGCAGGGAGCTCCGGGA CGCCGCCTCCCCATCCCCACGCTCGCATCAAGCTGGAGAACCCGCTGGACTACGCCAGCG CCTGGGCGGCTGCGGCGCGCAGTGCCGCTATGGGGACCTGGCGAGCCTGCATGGCGCGG CTGTAGCCCCCTACGCTACACTCGGCCCCCTCAGGGGCTGGCGGGCCAGGAAAGCGACT TCACCGCACCTGATGTGTGTACCCTGGCGGCATGGTGAGCAGAGTGCCCTATCCCAGTC ACATGCGTTTGGAGACTGCCAGGGACCATGTTTTGCCCATTGACTATTACTTTCCACCCC 2462 2522 1009 2582 1069 1129 2702 1189 1249 1309 2642 2762 2822 2882 2942 3002 1429 1489 3062 1549 3122 1609 3182 1669 3242 1729 3302 1789 1849 3482 3362 1969 δy g δý g οy g Óλ qq Ω Q δ οp Qy Db Qγ g Qy Db QQ Οÿ QQ qq qq ŏλ δy óλ qq Dp φ δ pp Qγ g δ g á

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The invention relates to isolated polynucleotide (I) and polypeptide (II) sequences. (I) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chromosome and gene mapping, and in recombinant production of (II). The 3408 3468 5041 3168 3228 4801 3288 4861 4921 4981 S 3469 GCCAGCCACACAAACGTTTACTTATCTTATGCCACGGGAAGTTTAGAGAGCTAAGATTAT nutations GCCAGCCACACAAACGTTTACTTATCCCACGGGAAGTTTAGAGAGCTAAGATTAT ICTGTGATGATCCTCATATGGCCCAGTGTCAAGTTGTGCTTGTTTACAGCACTACTGT 3409 TCTGTGATGATCCTCATATGGCCCAGTGTCAAGTTGTGCTTGTTTACAGCACTACTGT 3109 TCCACACCCAGTGAAGCATTGGAAACCCTATTTCCCCACCCCAGCTCATGCCCCTTTCA TCTCTTTCTCTTTTTTTTTTTTTTTTTTAACCCTCCCANGGCACCTTC CTATTGATGTACAGTCTGTCATGAACATGTTCCTGAATTCTATTTGCTGGGCTTTTTTT New isolated polynucleotide and encoded polypeptides, useful in diagnostics, forensics, gene mapping, identification of mutation responsible for genetic disorders or other traits and to assess biodiversity Human; chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; 5042 CIGGGGAAATCAAAACAAAAAACAAGCAAACAAAAAAAAA 5082 DNA encoding novel human diagnostic protein #29276. Claim 1; SEQ ID No 29276; 103pp; English. BP. RESULT 7 AAS93472/c ID AAS93472 standard; cDNA; 3590 Tang YT; 31-MAR-2000; 2000US-0540217. 23-AUG-2000; 2000US-0649167. 30-MAR-2001; 2001WO-US08631 (first entry) Drmanac RT, Liu C, WPI; 2001-639362/73. P-PSDB; ABG29285. (HYSE-) HYSEQ INC WO200175067-A2. Homo sapiens 13-FEB-2002 11-OCT-2001. AAS93472; 3529 4922 4982 4682 3169 4742 3229 4802 3289 g qq g ρp δ q ŏ δ qq qq δy δy

Polymucieculdes are also used in diagnostics as expressed sequence cays for identifying expressed genes. (1) is useful in gene therapy techniques to restore normal activity of [II] or to treat disease states involving quantitating a polypeptide in tissue, as molecular weight markers and as a food supplement. (II) and its binding partners are useful in medical imaging of sites expressing (II). (1) and (II) are useful for treating or disorders involving aberrant protein expression or biological activity. The polypeptide and polyncleotide sequences have applications in diagnostics, forensics, gene mapping, identification of mutations responsible for genetic disorders or other traits to assess biodiversity and to produce other types of data and products dependent on DNA and diagnostic coding sequences of data and protects dependent on DNA and diagnostic coding sequences of the invention.

Note: The sequence data for this patent did not appear in the printed security from WIPO usèd in diagnostics as expressed Sequence 3590 BP; 807 A; 977 C; 1016 G; 790 T; 0 other; ftp.wipo.int/pub/published_pct_sequences 855555555555555555555

1682 ITTGAGGCIGTCAGAGCGCTTTTIGCGTGGTTGCTCCCGCAAGTTTCCTTCTGGAGCT 1741 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1111 | 1 1502 TAATAACTCAGTTCTTATTTGCACCTACTTCAGTGGACACTGAATTTGGAAGGTGGAGGA 1561 1562 ITTTGTTTTTTTTTAAGAICTGGGCATCTTTTGAATCTACCCTTCAAGTATTAAGAG 1621 1622 ACAGACTGTGAGCCTAGCAGGGCAGATCTTGTCCACCGTGTGTCTTCTTCTGCACGAGAC 1681 1742 TCCCGCAGGTGGGCAGCTAGCTGCAGCGACTACCGCATCATCACAGGCCTGTTGAACTCTT 1801 CTGAGCAAGAAGGGGAGGCGGGGTAAGGGAAGTAGGTGGAAGATTCAGCCAAGCTCAA 1861 GAGGAGCTTTCCAGAATCTGTTCCAGAGCGTGCGGAAGTGATCCAGAACCCGGGCCCCA 1981 3170 GAGGAGCTITCCAGAAICIGTICCAGAGCGTGCGCAGAAGTGTATCCAGAACCCGGGCCCCA 3111 GGCACCCAGAGGCCGCGAGCGCACCTCCCGGCGCCAGTTTGCTGCTGCTGCAGCAGC 2041 2101 2102 AGCAGCAAGAGACTAGCCCCAGGCAGCAGCAGCAGCAGGAGGTGAGGATGGTTCTCCCC 2161 AAGCCCATCGTAGAGGCCCCACAGGCTACCTGGTCCTGGATGAGGAACAGCAACCTTCAC 2221 2933 AAGCCCATCGTAGAGGCCCCACAGGCTACCTGGTCCTGGATGAGGAACAGCAACCTTCAC 2874 2222 AGCCGCAGTCGGCCCTGGAGTGCCACCCCGAGAGAGGGTTGCGTCCCAGAGCCTGGAGCCG 2281 25; Gaps Score 3498.2; DB 23; Length 3590; Pred. No. 0; 0; Mismatches 3; Indels 25; 3; Indels 68.8%; 99.2%; Best Local Similarity 99.2 Matches 3565; Conservative Query Match 1802 1982 2993 2042 1922 2162 ò 셤 ò g ö qq δŽ g δ 셤 δŏ qq οy q g ŏ δ QQ οy a δλ g Ω

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                                                                                                                transforming
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                                                                                                                                                                                                                                                                                    Length 4180;
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                                                                                                                                                                                Complementary DNA sequences derived from the cDNA may be used detect the presence of androgen receptor (AR) mRNA in tumour to detect AR gene defects using DNA hybrisidation assays.
                                                                                                                                                                                                                                           Sequence 4180 BP; 1024 A; 1149 C; 1083 G; 924 T; 0 other;
                                                                                                                                                                                                                                                                                                              664; Indels
                                                                                                                                                                                                                                                                         Query Match
48.9%; Score 2486.4; DB 10;
Best Local Similarity 77.9%; Pred. No. 0;
Matches 3416; Conservative 0; Mismatches 664; In
                                Joseph DR, Lubahn
   NORTH CAROLINA
                                                                                                                                                        English
                                                                                                                                                      5; 41pp;
                                EW,
UNIVERSITY OF
                                                       WPI; 1989-324206/44
                               Wilson
                                                                           P-PSDB; AAP93110
                                                                                                                                                    Disclosure; Fig.
                               French FS,
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•	2648 3641 2708 3701	9 cdctallactrcccacccanadacctdcctdarctdtdcardadactrctgctrg 2 rectargedaccrcrcaardagacctgcaaggrctrctrcaardagaccgcrcaardgg 2 rectargedaccrcrcaargtggaagctgcaaggrctrctrcaardagaccgcrgaagg 1 1 1 1 1 1 1 1 1 1	258 358 264 364	oy Oy Oy
	3461 2528 3521 2588	CAGAGTGCCCTATCCCAGTCCCACTTGTGTCAAAAGCGAAATGGGCCCCTGGATGGA	340 246 346 252	oy oy oy
	3341 2408 3401 2468	2 CGGCGGCGGCGAGGCG 	328; 234; 334 240	0y 0y 0b
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•	3161 2273 3221 2327	2 GGGGAGCCTGCATGGCGGGGGGCGCGGGTCTCTGGGTCACCCTCAGCCGCCGCGCGCTIIIIIIIIII	310: 221. 316. 227.	67 67 67 67
	0 7	2 CCCGCTGGACTACGGCAGCGCCTGGGCGGCGGCGCGCAGTGCCGCTATGGGGACCT	3043	O.Y DD
	3041 2153	2 GGCGGACCGCCCCCCCCCCCCCCCCCCCCCCCCCCCCC	2983	QQ Op
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	2861 1973 2921 2033	TITCAAGGGGGGTTACACCAAGGGCTAGAGGCGGAGACCTAGCTGGTTGGCAGCGCTGTTTCAAGGGTTACACCTAGTTGGCAGCGCTAGTTGTAGTTGTAGGGTGAGAGTTGGGCTGGCT	2802 1914 2862 1974	27 27 27 27
× •	2/41 1853 2801 1913	ACTTTGGGAGTTCACCGCTGTGCTCCCACTCGTGTGCCCATTGCCCA GICTCTGGGAGTCCACCGCCGTGCTCCACTCCTTGTGCGCCCTTGGCCGA AGGTTCTCTGCTAGACGACAGCGCGAGCACACTCCTTGTGCGCCCTTGGCTA AGGTTCTCTGCTAGACGACAGCGCAGGCAAGACACTGAAGATACTGCTGAGTA III I IIIIIIIIIIIIIIIIIIIIIIIIIIIIIII	2682 1794 2742 1854	24
	68	ALCIGACACIGCCARSSASISISISIAAASCATCIGGGGGGGATTGCATGAGCCCCCGGGGGGATTGCATGTACGCCCCCGGGGTTGGAGGTTGCATGTACGCCCCGGGGTTGGAGGTTGCATGTACGCCCCAGGGGGGGG	73	9 2 9
	2621 1733	CATTICTGACAAGGCCAAGGAGTIGTGTAAGGCAGTGTCGGTGTGCCATGGGCCTGGGTGT	2562	ξ, q

4719 4778 4421 4481 3548 3668 3728 3188 4181 3248 4241 4301 3368 4361 3428 3488 4541 3608 4601 4661 4061 3068 3128 2768 3761 3821 2888 3881 2948 3941 3008 4001 4121 GCAGTGCCTT-GGGGAATTTCCTCTATTGATGTACAGTCTGTCATGAACATGTTCCTGAA GAAAGTCAAGCCCATCTATTTCCACACCCAGTGAAGCATTGGAAACCCTATTTCCCCACC TITCAATGAGTACCGCATGCACAAGTCCCGGATGTACAGCCAGTGTGTCCGAATGAGGCA CACATCCTGCTCAAGACGCTTCTACCAGCTCACCAAGCTCCTGGACTCCGTGCAGCCTAT CAGCCCCACTGAGGAGACACCCAGAAGCTGACACATGAACATTGAAGGCTNTGAATG CGTGGACGACCAGATGGCTGTCATTCAGTACTCCTGGATGGGGCTCATGGTGTTTGCCAT GGGCTGGCGATCCTTCACCAATGTCAACTCCAGGATGCTCTACTTCGCCCCTGATCTGGT 3609 4720 4542 4602 3669 4662 3729 4422 3489 4062 4122 3189 4182 4242 3309 4302 3369 2709 3882 2949 3942 3009 4002 3069 2769 2829 3822 2889 3702 3762 Dβ g δy g ζŎ g ŏ g QΥ g δy 요 Q q QΥ qq δλ qq QΥ g δλ qq Q. οy qq Db QΫ g δy g Qγ g QY QY δλ

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4958
                                                                                                                                                                                                                                                  Human; ds; astrocytoma; cytostatic; staging; cysteine methylation; CpG;
bisulphite; brain tissue; MALDI; ESI; electron spray mass spectrometry;
matrix assisted laser desorption/ionization mass spectrometry.
                     TCTAACCCICCCAIGGCACCTICAGACTITGCTTCCCATTGIGGCTCCTAICTGTGTTTT 4898
                                                3788 GCATIGCCTIGGGGGAAATICCTCTACTGATGIACAGICIGICATGAACAIGTICCCCAA 3847
                                                                                                   GAATGGTGTTGTATGCCTTTAAATCTGTGATGGTCCTCATATGGCCCCAGTGTCAAGTTGT
                                                                         Human DNA for staging of Astrocytomas, complement, #50.
                                                                                                                                                                                                                                                                                                                                                                     Berlin K;
                                                                                                                                                                                              ABK34013 standard; DNA; 3715 BP.
                                                                                                                                                                                                                                                                                                                     02-JUL-2001; 2001WO-EP07538
                                                                                                                                                                                                                                                                                                                                 30-JUN-2000; 2000DE-1032529.
01-SEP-2000; 2000DE-1043826.
                                                                                                                                                                                                                          (first entry)
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                                                                                                                                                                                                                                                                                                                                                      (EPIG-) EPIGENOMICS AG.
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methylation states of the CpG dinucleotides of (I). The array is useful CfG for determining genetic and/or epigenetic parameters, classification, differentiation, grading, staging, treatment and/or diagnosis of astrocytomas, or the predisposition to astrocytomas by analysing cytosine bases which are comethylations, involves obtaining a biological sample containing genomic DNA, extracting the genomic DNA, converting cytosine bases which are unmethylated at the 5-position, in the genomic DNA sample, to uracil or behaviour, by chemical treatment and amplifying chemically pre-treated genomic DNA fragments using the array and a polymerase, where the amplificates carry a detectable label. The method further involves defecting methylation status of one or more cytosine positions, and analysing methylation status of one or more cytosine positions, and analysing methylation status of the cytosine positions by reference to bisulphite, hydrogen sulphite or disulphite. The amplificates carry a challysing methylation status of the cytosine position or status of brain tissue, based on the specific genomic methylation status of brain tissue, based on the specific genomic methylation status of brain tissue, based on the specific genomic methylation status of brain tissue, based on the specific genomic methylation status of brain tissue, based on the specific genomic methylation status of content and the specific genomic methylation status of content and the specific genomic methylation status of brain tissue, based on the specific genomic methylation status of content and the specific genomic methylation status of content and the specific genomic sequence of chemically, the maps spectrometer. The fragments of chemically care detected has mass spectrometer. The fragments of chemically contents assisted laser described have a single positive or regarded by matrix assisted laser described expection spray mass spectrometry ference bush
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          1341 CGGAGAGAACCTTGTTTCCCCACTCTCTCTCCACCTCCTCCTGCCTTCCCCACCC 1400
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Pred. No. 0:
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Pred. No. 0;
0; Mismatches 848; Indels
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Sequence 3715 BP; 818 A; 149 C; 1055 G; 1693 T; 0 other;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           ftp.wipo.int/pub/published_pct_sequences.
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76.18;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           samples of the invention.
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Matches 2838; Conserv
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The invention relates to a nucleic acid comprising a sequence (I) of at least 18 bases in length of a segment of chemically pre-treated genomic DNA which has any one of the sequences of (ABK33919-ABK34032) or its complement. Also included are an Oligonucleotide or peptide nucleic acid (or set thereof) of at least 9 nucleotides which hybridises to (I), probes for detecting cytosine methylation or singlenucleotide polymorphisms (SNP) in (I), an array of oligomers or peptide nucleic acids for analysing diseases associated with the

Novel chemically modified genomic DNA sequences, useful in the characterisation, classification, differentiation, grading, staging, treatment and/or diagnosis of astrocytomas or predisposition to

Claim 1; SEQ ID No 100; 37pp; English.

astrocytomas

1032 3791 1272 1212 3911 1152 3971 1452 3671 3731 1692 1632 3491 1572 1512 3611 1872 1812 3311 1752 3371 3431 3551 3080 1992 3200 3251 CCTGATCTGTGGAGATGAAGCTTCTGGGTGTCACTATGGAGCTCTCACATGTGGAAGCTG TTATGAAGCAGGGATGACTCTGGGAGCCCGGAAGCTGAAGAAACTTGGTAATCTGAAACT 3492 GGAGACTGCCAGGGACCATGTTTTGCCCATTGACTATTACTTTCCACCCCAGAAGACCTG CAAGGICTICTICAAAAGAGCCGCTGAAGGGAAACAGAAGTACCTGTGCGCCAGCAGAAA ACAGGAGGAAGGAGGCTTCCAGCACCACCAGCCCCACTGAGGAGACAACCCAGAAGCT CTACGGCTACACTCGGCCCCTCAGGGGCTGGCGGCCAGGAAAGCGACTTCACCGCACC 1691 TAATATATAATACCCTAACGACATAATAAACAAAATACCCTATCCCAATCCCAATTATAT CAAAAGCGAAATGGGCCCCTGGATGGATAGCTACTCCGGACCTTACGGGGACATGTTT 3141 TGGGTCACCTCAGCCGCCGCTTCCTCATCCTGGCACACTCTTCTCACAGCCGAAGAAGG 1931 TAAATCACCCTCAACGCCGCTTCCTCATCCTAACACACCTCTTCACAACCGAAAAAA CCAGTTGTATGGACCGTGTGGTGGTGGTGGGGGGT------GGTGGCGGCGGCGGCGG 1871 CCAATTATATAAACCGTATAATAATAATAAAAATAATAATAATAACGACGACGACGACGACGA TGATGTGTGGTACCCTGGCGGCATGGTGAGCAGAGTGCCCTATCCCAGTCCCACTTGTGT CTACAACTTTCCACTGGCTCTGGCCGGACGCCGCCCCCTCCGCCGCCTCCCCATCCCCA GCAGTGCCGCTATGGGGACCTGGCGAGCCTGCATGGCGGGGTGCAGCGGGACCCGGTTC CGCTCGCATCAAGCTGGAGAACCCGCTGGACTACGGCAGCGCCTGGGCGGCTGCGGCGGC 1091 1331 1271 3912 1151 3612 3672 1391 3792 3972 3732 3552 3081 3252 3312 1751 3372 3432 1631 3201 3021 g QQ qq g g ò δy qq q δ ΩD $Q\underline{Y}$ pp δ ΩD QY $\dot{\Omega}$ δŏ qa :: qq qq δ οy δy qq QΥ QQ δŽ QΥ QQ Qγ QQ QΥ g QΥ

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CTCCTGGATGGGGCTCATGGTTTTGCCATGGGCTGGCGATCCTTCACCAATGTCAACTC 4151
                                                           CAGGATGCTCTACTTCGCCCCTGATCTGGTTTTCAATGAGTACCGCATGCACAAGTCCCG 4211
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                                                                                                                      CACCCCCAGGAATTCCTGTGCATGAAAGCACTGCTACTTCTAGCATTATTCCAGTGGA 4331
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                                                                    4632 GIGAAGCAIIGGAAACCCIAIIICCCCACCCCAGCICAIGCCCCCIIICAGAIGICIIC 4691
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        CAAGGCCTTGCCTGGCTTCCGCAACTTACACGTGGACGACCAGATGGCTGTCATTCAGTA
                                      GATGTACAGCCAGTGTGCCGAATGAGGCACCTCTCTCAAGAGTTTGGATGGCTCCAAAT
                                                                                                                               TGGGCTGAAAAATCAAAAATTCTTTGATGAACTTCGAATGAACTACATCAAGGAACTCGA
                                                                                                                                                            CACCAAGCTCCTGGACTCCGTGCAGCCTATTGCGAGAGCTGCTTCAGTTCACTTTTGA
                                                                                                                                                                                                                        CTCTGTGCAAGTGCCCAAGATCCTTTCTGGGAAAGTCAAGCCCATCTATTTCCACACCCA
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The present invention relates to chemically modified DNA sequences of signal transduction associated genes. The DNA sequences are chemically modified using a solution of bisulphite, hydrogen sulphite or modified using a solution of bisulphite, hydrogen sulphite or dislushments. Also disclosed are oligonucleotides and/or PNA oligomers for detecting the cytosine methylation state (CpG islan's) of these genes, and a method for the diagnosis and/or therapy of genetic and epigenetic parameters of genes associated with signal transduction. The genomic DNA can be obtained from cells or cellular components which contain DNA, e.g. cell lines, biopsies, blood sputum, stool, urine, cerebral-spinal fluid, tissue embedded in paraffin such as tissue from eyes, intestine, kidney, brain, heart, prostate, lung, breast or liver, histologic object slides, and all their possible combinations. The sequences of the invention are useful for the diagnosis and therapy of diseases associated with signal transduction, or sequences of different genes associated with signal transduction, or the component of the invention are useful for the signal transduction, or the component of different genes associated with signal transduction, or
                                                                                                                                                                                    Human, signal transduction associated gene; cytosine methylation state; CpG island; signal transduction associated disease; solid tumour; cancer; antitumour; cytostatic; mutant; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            form part of the printed
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        Oligonucleotide for diagnosis and therapy of diseases associated with signal transduction e.g. cancer, comprises chemically modified genomic sequences of genes associated with signal transduction
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                their complementary sequences. Note: The sequence data for this patent did not form part of the printe specification, but was obtained in electronic format directly from the European Patent Office.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           1401 GAGTGCGGAGCCAGAGATCAAAAGATGAAAAGGCAGTCAGGTCTTCAGTAGCCAAAAAAC 1460
                                                                                                                                                      gene modified complementary DNA #194.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     44.2%; Score 2244.2; 76.1%; Pred. No. 0; ive 0; Mismatches
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 Claim 1; SEQ ID No 388; 24pp; English.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Berlin K;
                                                                                                                                                    Signal transduction associated
                                        DNA; 3715
                                                                                                                                                                                                                                                                                                                                                                                                                                   30-JUN-2000; 2000DE-1032529.
01-SEP-2000; 2000DE-1043826.
                                                                                                                                                                                                                                                                                                                                                                                             29-JUN-2001; 2001WO-EP07472.
                                                                                                              (first entry)
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Matches 2838; Conservative
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            (EPIG-) EPIGENOMICS AG
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                                      ABK31545 standard;
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                                                                                                                                                                                                                                                                                       Synthetic.
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1461 AAAACAAACAAAAAAAGAAAAAAAAAAAAAAAAAAAAA	GGGCAGATCTTGTCCACCGTGTGTCTTCTTCTGCACGAGACTTTGAGGCTGTCAGAGCGC		3107 ATTCCAAACGCGCAAATATTCCAAACCCCCAAACCCCCAAAAACGGGAA 2001 CGCAGCACCTCCGGCGCAGTTTGCTGCTGCTGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAG	29121 CAGGCAGCAGCAGCAGGGTGAGGATCTCCCCCAAGCCCATCGTAGAGGCCCCCCCC	2301 GCTGCCGCAGCAGCACCTCCGGACGAGGATGACTCAGCTGCCCATCCACTT 1 1 1 1 1 1 1 1 1 1
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3140 1992 1572 3611 3671 3080 1812 3311 1752 3431 3491 3551 2900 2960 2112 3020 2052 2412 2780 2292 2840 2232 2172 CGACGACGACGACGACGACGACGACGACGACGACGAAACGAAAACTATAAACCC 3432 CAAAAGCGAAATGGGCCCCTGGATGGATAGCTACTCCGGACCTTACGGGGACATGCGTTT 3492 GGAGACTGCCCAGGGACCATGTTTGCCCATTGACTATTACTTTCCACCCCAGAAGACCTG CCTGATCTGTGGAGATGAAGCTTCTGGGTGTCACTATGGAGCTCTCACATGTGGAAGCTG CGCTCGCATCAAGCTGGAACCCGCTGGACTACGGCAGCGCCTGGGCGGCTGCGGCGGC CCAGITGIAIGGACCGIGIGGIGGIGGIGGIGGIG------GGIGGCGCGCGGCGGCGG 3372 TGAIGIGIGGIACCCIGGCGCAIGGIGAGCAGAGIGCCCIAICCCAGICCCACIIGIGI 2051 CGCTCGCATCAAACTAAAAAACCGCTAAACTACGACAAACGCCTAAACGACTACGACGAC GCAGTGCCGCTATGGGGACCTGCGAGCCTGCATGGCGCGGGTGCAGCGGGACCCGGTTC CCTAGGCTGCTCTGGCAGCGCTGCAGCAGGAGCTCCGGGACACTTGAACTGCGGTCTAC CCTGTCTCTACAAGTCCGGAGCACTGGACGAGGCAGCTGCGTACCAGAGTCGCGACTA GGTGTCCATGGGCCTGGGTGTGGAGGCGTTGGAGCATCTGAGTCCAGGGGAACAGCTTCG GGGGGATTGCATGTACGCCCCACTTTTGGGAGTTCCACCCGCTGTGCGTCCCACTCCTTG AAAAAATTACATATACGCCCCCCTTTTAAAATTCCACCCGCTATACGTCCCACTTA 1571 3552 3612 1811 3141 1931 3201 3252 1991 2411 2291 2961 3021 3081 2661 2351 2781 2841 2231 2901 2171 2111 2601 2721 qа . q QQ g ΟŽ Op ò ŏ QQ Ω 염 q Qγ QΥ Qγ QQ g δ g δŽ g δ QΩ οy q Οŷ οy Q QΫ́ Ω οy q δŽ Ω

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qq	1451	CAAAATCTTCTTCAAAAAAACCGCTAAAAAAAAAAAAAA	
ŏλ	3672		0y 47
QQ	1391	TAATTACACTATTAATAAATTCCGAAAAAATTATCCATCTTATCGTCTTCGAAAATA 1332	Dp 3
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Д	1331	TTATAAAACAAAAATAACTCTAAAAACCCGAAAACTAAAAAACTTAATAATCTAAAACT 1272	Dp 3
Q Q	3792	ACAGGAGGAAGGAGGCTTCCAGCACCACCAGCCCCACTGAGGAGACAACCCAGAAGCT 385	Oy 48
٥	3852	GACAGTGTCACACACCACCACACCACCACCACCACCACCACACCCCAAAA	Qy 49
² 연	1211	ANGEST OF THE STATE OF THE STAT	Dp 1
δλ	3912	CATTGAGCCAGGTGTAGTGTGTGCTGGACACAACAACAACCAGCCCGACTCCTTTGCAGC 39	Qy 49
QQ	1151	CATTAPACCAAATATAATATATATACTAAACAACCAACCCGACTCCTTTACAAC 1092	qa
Qy Db	3972	CTTGCTCTCTAGCCTCAATGAACTGGGAGAGACAGCTTGTACACGTGGTCAAGTGGGC 4031 	Qy 50.
Oy Db	4032		RESULT AAQ1200
δy	4092		XX XX
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Oy Dp	4152		XX DE Fu
Qy Db	4212		XX Kai
QY	4272	CACCCCCCAGGAATTCCTGTGCATGAAAGCACTGCTACTTCTACAGCATTATTCCAGTGGA 4331	FT CD
qq	791		FT
Oy Db	4332	TGGGCTGAAAAATCAAAAATTCTTTGATGAACTTCGAATGAACTACATCAAGGAACTCGA 4391 	XX PN WOS XX PD 30
0 <u>7</u>	4392	TCGTATCATTGCATGCAAAAGAAAAATCCCACATCCTGCTCAAGACGCTTCTACCAGGT 4451 	XX PF 19- XX DP 17-
Qy Db	4452		XX XX XX (AF XX
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Qy Db	371	47 31	CC DNA CC there

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872 TCCCATTGTGGCTCCTATCTGTGTTTTGAATGGTGTTGTATGCCTTTAAATCTGTGATGA 4931
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                                                                                                                                                                                    us sequence was isolated by screening a rat ventral prostate numbda gtll library in E.coli Y1090. Initial screening.

We with probes designed for homology to nucleotide sequences in the Ab-binding domain of known steroid receptors. Positive clones were nen screened with 24mer probes specific for the various steroid seceptors to eliminate those which coded for known receptors. Any amaining clones were analysed by restriction mapping and were
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Location/Qualifiers
33..274
/*tag=
/*toduct= full-length (902 amino acids) rAx
/product= full-length ras residue shorter rAR protein"
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               111-length rat androgen receptor coding sequence.
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AQ12002 standard; DNA; 3217 BP.
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22 CAAAACAAAA 12
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PSDB; AAR12224.
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11; 2428 2451 2617 2188 2248 2308 2368 2497 2557 CCCCACTITIGGGAGTICCACCCGCTGTGCGTCCCACTCCTTGTGCCCCATGGCCGAAT 2737 2011 CCGGCGCCAGITIGCIGCIGCIGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGC 2128 GTGTGGAGGCGTTGGAGCATCTGAGTCCAGGGGAACAGCTTCGGGGGGATTGCATGTACG 2677 GCAAAGGTTCTCTGCTAGACGACAGCGCAGGCAAGAGCACTGAAGATACTGCTGAGTATT 2797 TCTACCCTCGGCCGCCGTCCAAGACCTACCGAGGAGCTTTCCAGAATCTGTTCCAGAGCG 1951 768 468 528 588 648 708 348 408 1891 228 9 Gaps ACCTGGTCCTGGATGAGGAACAGCAACCTTCACAGCCGCGGTCGGCCTGGAGTGCCCACT AGCAGCAGCAGCAGCAACAGCAGCAGGAGGTAATATCCGAAGGCAGCAGCAGCAGA GAGCGAGGGAGCCTCGGGGGCTCCCACTTCCTCCAAGGACAATTACTTAGGGGGCACTT 649 GAGCAAGGGAGCCACTGGGGGCTCCCTTCCTCCAAGGATAGTTACCTAGGGGGCAATT CGACCATTTCTGACAACGCCAAGGAGTTGTGTAAGGCAGTGTCGGTGTCCATGGGCCTGG AGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAAGAGACTAGCCCCA---GGCAGC CTGAGAGCGCTGCCTCCCCGGAGCTGCTGCTGCTCCTGGCAGGGCTCCTGCCGC AGCAGCTGCCAGCACCTCCGGACGAGGATGACTCAGCTGCCCCATCCACGTTGTCCCTGC TGGGCCCCCACTTTCCCCCGGCTTAAGCAGCTGCTCCGCTGACCTTAAAGACATCCTGAGCG AGCAGCAGCAGCAGGTGAGGATGGTTCTCCCCAAGCCCCATCGTAGAGGCCCCCACAGGCT CCGAGAGAGGTTGCGTCCCAGAGCCTGGAGCCGCCGTGGCCGCCAGGAAGGGGCTGCCGC GAAGTAGGTGGAAGATTCAGCCAAGCTCAAGGATGGAAGTGCAGTTAGGGCTGGGAAGGG Indels Sequence 3217 BP; 776 A; 873 C; 843 G; 725 T; 0 other; is given here. DB 12; 393; Score 2189.2; Pred. No. 0; 0; Mismatches rat AR coding sequence ;0 AGGCCAGCACCATGCAACTCCTT-43.1%; 82.4%; 181 CCGGTGCCTGTTT-----Conservative Best_Local Similarity Matches 2726; Conserv The sequenced. Match 692 349 2452 589 2498 2618 2678 2738 2129 2189 2249 2309 409 2369 2429 529 709 829 289 469 1952 2012 2072 1832 1892 61 121 194 229 Query g ρp Ω g δy pp ò Бb g g οy δ ΟÝ 임 ΟŸ Dp οy οqα δý δý δŽ g δλ g οy qq QΫ́ q

1743 3337 3577 3637 3877 1188 3157 3217 3397 3457 3037 2857 2977 1624 CCATCGACTATTACTTCCCACCAGAAGACCTGCCTGATCTGTGGAGATGAAGCTTCTG CCACCAGCCCCACTGAGGAGACAACCCAGAAGCTGACAGTGTCACACATTGAAGGCTATG ATAGCTACTCCGGACCTTACGGGGACATGCGTTTGGAGACTGCCAGGGACCATGTTTGC CCATTGACTATTACTTTCCACCCCAGAAGACCTGCCTGATCTGTGGAGATCAACCTTCTG **AAGGGAAACAGAAGTACCTGTGCGCCAGCAGAAATGATTGCACTATTGATAAATTCCGAA** GCGGCGGCGGCGGGGGGGGGGGGTGTAGCCCCTACGGCTACACTCGGCCCCCTCAGG CCGGAGCACTGGACGAGCTGCGTACCAGAGTCGCGACTACTACAACTTTCCACTGG AGAACCCGCTGGACTACGGCAGCGCCTGGGCGGCGGCGGCGCGCAGTGCCGCTATGGGG ACCTGGCGAGCCTGCATGGCGCGGGTGCAGCGGGACCCGGTTCTGGGTCACCCTCAGCCG CCCCTTTCAAGGGAGGTTACACCAAAGGGCTAGAAGGCGAGAGCCTAGGCTGCTCTGGCA GCGCTGCAGCAGCAGCACCTCCGGGACCACTTGAACTGCCGTCTACCCTGTCTCTACAAGT 3638 1744 3698 1864 3818 1564 1804 3758 3338 3458 3518 2918 1069 2978 1129 3038 1189 3098 1249 3158 1309 3218 1367 3278 1384 2858 1009 2798 949 889 qq q q δ g Ω g οy В δy 8 Qγ g δy Pp ò Db δ d q δ Q Óγ g Ŏλ qq δ g óλ g δy g δy qq Ω

Fri May

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4955 ITGTGCTTGTTTACAGCAC--TACTCTGTGCCAGCCACAAACGTTTACTTATG
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/*tag= a

/product=98 kD polypeptide

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/*tag= b

/product=79 kD polypeptide
                                                                                                                                                                                                                            Location/Qualifiers
                                                                                                                                                                                          Androgen receptor; TR2 polypeptide;
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                                                                                                                                AAN91578 standard; DNA; 3217
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Matches 2715; Conservative
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P-PSDB; AAP91006.
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              AATGTCAGCCCATCTTTCTGAATGTCCTGGAAGCCATTGAGCCAGGTGTAGTGTGTGCTG
                       GACACGACAACAACCAGCCCGACTCCTTTGCAGCCTTGCTCTAGCCTCAATGAACTGG
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             New DNA encoding new androgen receptor and TR2 polypeptide(s) - abito bind DNA, and derived antibodies, useful for receptor assay and
                                                                                                                                                                                          The sequence is used to express the corresp. peptides and for hybridisation assays of RNA and DNA encoding androgen receptors. The 98 kD product starts at the first Met codon; the 79 kD product starts from the second
                                                                                                                                                                                                                                                                                                                                                                                                                        Length 3217;
                                                                                                                                                                                                                                                                                                                                                                                                       43.0%; Score 2185.8; DB 10; Length 3217 82.1%; Pred. No. 0; Live 0; Mismatches 407; Indels 185;
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 ATCG	CCAG	AGCA	AGC?	7000	CAC) - -	SACT	16CT	CCAGCACCATGCAACTCCTT	AACAACAGCAGCAGGAAGCAGTATCCGAAGGCAGCAGCAGCGGAGAGAGA	CAGGGAGCCTCGGGGGCTCCCACTTCCTCCAAGGACAATTACTTAGGGGCACTTCGA [CCATTICIGACACGCCAAGGAGTIGTAAGGCAGTGTCGGTGTCCAIGGCCTGGGTC 	TGGAGGCGTTGGAGCATCTGAGTCCAGGGGAACAGCTTCGGGGGGATTGCÀTGTACGCCC 	CACITITIGGAGITCCACCGCTGTGCGTCCCACTCCTTGTGCCCATTGGCCGAATGCA 	aaggitcicigciagacgacagcgcaggcaagacacigaagataciccgagiaticc 	CTTTCAAGGGGGGTTACACCAAAGGGCTAGAAGGCGAGAGCCTAGGCTGCTGGGGGGCGC 	CTGCAGCAGGAGCTCCGGGACACTTGAACTGCCGTCTACCTGTCTCTACAAGTCCG 	GAGCACTGGACGAGGCAGCGTACCAGAGTCGCGACTACTACAACTTTCCACTGG
ACCT	3966	CAGC	CAGC	TCTC	CCT	GGAC	GATC 	AGC.	3CAG	SCAG IIII SCAG	CACT CTCT	GTTG	TCCA	TGTG	- - - - - -	AGGC	ACT	GTA(
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1072 GAGCAGTAGACGAGCAGCATACCAGAATCGCGACTACTACAACTTTCCGCTCGCT	AGCTGGAGA 	041 ACCGGCTGGACTACGGCAGCGCCTGGGCGGCTGCGGCGCGCAGTGCCGCTATGGGACCCTATGGACTACGGCAGTACGGCGCGCGGCGGCGGCGCGCAGCGCAATGCGGACTATGGGGACT	101 TGGCGAGCCTGCATGCGGGTGCAGCGGGACCCGGTTCTGGGTCACCCTCAGCGCGCG	52 TGGCTAGCCTACATGGAGGGAGTGTAGCCGGAACCAGCACCAGCATGGATGCGGGGGGGG	11 1 1 1 1 1 1 1 1 1	221 GTGGTGGTGGGGGTGGTGGCGGCGGCGGCGGCGGCGGCGG	- INSTRUCTION OF THE PROPERTY	387 GIAGCCCAAGCGATGCTGGGCCTGTAGCCCTATGCTACACACTCGGCCCCTCT	GCGACTTCACCGCACCTGATGTGTGTGGTACCCTGGCGG	GCAAGCCAGGAGGTGACTTCTCTGCCTCTGAAGTGTGGTATCCTGGTGGAGTTGTG	40	CCCCTATICCCAGGCACATGCGTTTGGAGACTGCCAGGGACCATGTTTGCCC	26	521 TTGACTA	3ACTATTACTTCCCACCCCAGAAGACCTGCCTGATCTGTGGAGATGAAGCTTCTG	3581 GTCACTATGGAGCTCTCACATGTGAAGCTGCAAGGTCTTCTTCAAAAGGCGCGCVAAG 1687 [11] 1 1 1 1 1 1 1 1 1	641 GGAAACAGAAGTACCTGTGCGCCA		701 AAAATTGTCCATCTTGTCGTCTTCGGAAATGTTATGAAGCAGGGATGACTC 	61 GGAAGCTGAAGAAACTTGGTAATCTGAAACTACAGGAGGAAGGA	CAGE	3821 CCAGCCCACTGAGGAGACACCCAGAAGCTGACAGTGTCACACATTGAAGGCTATGAAT	GCCCCACTGAGGACCCATCCCAGAAGATGACTGTATCACACATTGAAGGCTATG	00	987 GTCAACCTATCTTTAATGTCCTGGAAGCCATTGAGCCAGGAGTGGTGGTGGAGGGGGGGG	3941 ACGACAACAACCAGCCGACTTTTGCAGCTTTCCTCTCTTTGCTCTTTTGCTCTTTTTTTT	001 AGAGACAGCTTGTACACGTGGTCAAGTGGGGCCAAGGCCTTGCCTGGCTTCC	2107 AGAGACAGCTTGTACATGTGGTCAAGTGGGCCAAGGCCTTGCCTTGCCTTCCGCAACTT
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ABK34012 standard;

RESULT 13 ABK34012 ABK34012;

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                               TGGGCTGGCGATCCTTCACCAATGTCAACTCCAGGATGCTCTACTTCGCCCCTGATCTGG
                                                               TTTTCAATGAGTACCGCATGCACAAGTCCCGGATGTACAGCCAGTGTGTCCGAATGAGGC
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Claim 1; SEQ ID No 99; 37pp; English.

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Human; ds; astrocytoma; cytostatic; staging; cysteine methyiation; CpG;
bisulphite; brain tissue; MALDI; ESI; electron spray mass spectrometry;
matrix assisted laser desorption/ionization mass spectrometry.
                                                                                                                                                                                                                   Novel chemically modified genomic DNA sequences, useful in the characterisation, classification, differentiation, grading, staging, treatment and/or diagnosis of astrocytomas or predisposition to
                                            Human DNA for staging of Astrocytomas #50.
                                                                                                                                                                                        Berlin K;
                                                                                                                                   02-JUL-2001; 2001WO-EP07538
                                                                                                                                                 30-JUN-2000; 2000DE-1032529.
01-SEP-2000; 2000DE-1043826.
                             (first entry)
                                                                                                                                                                                       Olek A, Piepenbrock C,
                                                                                                                                                                         (EPIG-) EPIGENOMICS AG.
                                                                                                                                                                                                     WPI; 2002-171649/22
                                                                                                      WO200202808-A2.
                                                                                        Homo sapiens.
                             18-JUN-2002
                                                                                                                                                                                                                                           astrocytomas
                                                                                                                     10-JAN-2002.
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The invention relates to a nucleic acid comprising a sequence (I) of at cc least 18 bases in length of a segment of chemically pre-treated genomic DNA which has any one of the sequences of (ARK33919-ARK3403) or its complement. Also included are an oligonuclectide or peptide nucleic CC acid (or set thereof) of at least 9 nucleotides which hybridises to (I), probes for detecting cytosine methylation or single-cc primers for (I), probes for analysing diseases associated with the nector peptide nucleic acids for analysing diseases associated with the nector peptide nucleic acids for analysing diseases associated with the cc or peptide nucleic acids for analysing diseases associated with the methylation states of the CpG dinucleotides of (I). The array is useful differentiation, grading, staging, treatment and/or diagnosis of astrocytomas, or the predisposition to astrocytomas by analysing cytosine methylations, unvolves obtaining a biological sample cortaining quencin commethylations, unvolves obtaining a biological sample cortaining quencin commethylations, unvolves obtaining a biological sample cortaining appropriated at the 5-position, in the genomic DNA sextacting the genomic DNA, converting cytosine bases, which are cannother base which is dissimilar to cytosine pass, increased amplificates carry a detectable label. The method further, involves and electable label. The method further, involves of elentifying methylation status of one or more cytosine positions by teference to analysing methylation status of hen cytosine positions by terefrance to bisulphite. Nydrogen sulphite or disulphite. The amplificates carry a detectable model or more cytosine positions by tester detected in a mass spectrometer. The fragments are detected in an ass spectrometer. The fragments of the mass spectrometer or paint tissue, based on the specific genomic methylation status of the mass spectrometer. The fragments are detected in a mass spectrometer. The fragments are detected in a mass spectrometer. The fragments are detected in a

3172 CAAAAAA 3178

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                                                                                                                                                                                                                                                                      AGGGCAGATCTTGTCCACCGTGTGTCTTCTTGTGCAGGACTTTGAGGCTGTCAGAGCG
                                                                                                                                                                                                                                         reference
                                                                           45;
                                                              Length 3715;
 present sequence is one of the chemically pre-treated referen samples of the invention.

Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at
                                                                            Indels
                                                 other;
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                                                               DB
                                                 841 A; 149 C; 1001 G; 1724
                                                             Score 2154; DB
Pred. No. 0;
0; Mismatches
                                    ftp.wipo.int/pub/published_pct_sequences
                                                               42.48;
74.58;
                                                                       Similarity 74.5
32; Conservative
                                                  Sequence 3715 BP;
                                                                              Matches 2782;
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4451 TCACCAA 4451 TCACCAA 4514 TTATTAA 4514 ACTGCT 4514 ACTGCT 451 ACTGAA 4531 AGTGAAC 4631 AGTGAAC 4631 AGTGAAC 4631 AGTGAAC 4631 TCCCTT 4751 TACAGTC 4811 CTCCTTT 4811 CTCCTTT 4811 CTCCTTT 4811 TTTTTAT 4811 TTTTTAT 4931 ATCCCAA 3573 ATTTTTAT 4931 ACACACCA 3573 ATTTTTAT 4931 ACACACCA 3633 ATTTTTAT 4931 ACCTCCAA 3633 ATTTTTAT 4931 ACCTCCAA 3633 ATTTTTAT 4931 ACCTCCAA 3633 ATTTTTAT 4931 ACCTCCAA 3633 ATTTTTAT 4931 ACCTCCAA 3633 ATTTTTAT 4931 ACCTCCAA 3633 ATTTTTAT	ABK31544 ID ABK31544 AC ABK31544; XX XX DT 23-APR-2002 XX XX XX XX XX XX XX XX XX
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1964 3371 2024 3431 2084 3431 2144 351 2204 3671 2384 3731 2384 3731 2384 3911 2564 4031	4031 2684 4091 2744 4151 2804 4271 2924 4331 2984

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al transduction associated gene; cytosine methylation state; signal transduction associated disease; solid tumour; cancer; cytostatic; mutant; ds.
                         AAGCTCCTGGACTCCGTGCAGCCTATTGCGAGAGAGCTGCATCAGTTCACTTTTG 4510
                                                                                                                             TAATCAAGTCACACATGGTGAGCGTGGACTTTCCGGAAATGATGGCAGAGATCA 4570
                                                                                          TTATTGTATGTAAAAAAAAATTTTATATTTTGTTTAAGACGTTTTATTAGT 3103
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2000DE-1043826.
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The present invention relates to chemically modified DNA sequences of signal transduction associated genes. The DNA sequences are chemically modified using a solution of bisulphite, hydrogen sulphite or disulphite. Also disclosed are oligonucleotides and/or PNA oligomers of disulphite. Also disclosed are oligonucleotides and/or PNA oligomers of detecting the cytosine methylation state (CpG islands) of these genes, and a method for the diagnosis and/or therapy of genetic and epigenetic parameters of genes associated with signal transduction. The genomic DNA can be obtained from cells or cellular components which contain DNA, e.g. cell lines, biopsies, blood, sputum, stool, urine, cerebral-spinal fluid, tissue embedded in paraffin such as tissue from estabral-spinal fluid, tissue embedded in paraffin such as tissue from itsologic object slides, and all their possible combinations. The sequences of the invention are useful for the diagnosis and therapy of diseases associated with signal transduction e.g. solid tumnours and sequences of different genes associated with signal transduction, or sequences of different genes associated with signal transduction, or their complementary sequences. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from the Oligonucleotide for diagnosis and therapy of diseases associated with signal transduction e.g. cancer, comprises chemically modified genomic sequences of genes associated with signal transduction $\,$ Claim 1; SEQ ID No 387; 24pp; English χ; Piepenbrock C, European Patent Office. EPIGENOMICS WPI; 2002-147896/19 olek A,

Sequence 3715 BP; 841 A; 149 C; 1001 G; 1724 T; 0 other;

1639 1699 AGCTGCAGCGACTACCGCATCATCACAGCCTGTTGAACTCTTCTGAGCAAGAGAAGGGGGA 1819 GGCGGGGTAAGGGAAGTGCGAAGATTCAGCCAAGCTCAAGGATGGAAGTGCAGTTAG 1879 1340 GCGGAGAGAACCCTCTGTTTTCCCCCACTCTCTCTCCTCCTCCTCCTGCCTTCCCCACC 1399 1459 1700 CITITIGCGIGGIIGCICCCCCCAAGIIICCIICICIGGAGCIICCCGCAGGIGGGCAGCI 1759 247 367 427 187 68 CGAGTGCGGAGCCAGAGATCAAAAGATGAAAAGGCAGTCAGGTCTTCAGTAGCCAAAAAA AGGGCAGATCTTGTCCACCGTGTGTCTTCTTGTACACGAGACTTTGAGGCTGTCAGAGCG TITITICCGTGGTTGTTTTCGTAAGTTTTTTTTTTTTTGGAGTTTTTTCGTAGGTGGGTAGTTT 45; Query Match
Best Local Similarity 74.5%; Pred. No. 0;
Matches 2782; Conservative 0; Mismatches 905; Indels 45; 1640 1760 69 1520 248 308 368 1820 1400 188 1580 428 qq g ò g qq ò ò g οy q δ g οy ò ğ ă 셤

2779 2839 1243 2659 2239 1999 2059 883 667 GGCGGGGTAAGGGAAGTAGGTGGAAGATTTAGTTTAAGGTTAAGGATGGAAGTGTAGTTAG 547 607 GCCTAGGCTGCTGGCAGCGCTGCAGGAGCTCCGGGACACTTGAACTGCGGTCTA 2720 GIGCCCCAIIGGCCGAAIGCAAAGGIICTCTGCIAGACGACAGCGCAGGCAAGAACACIG AAGATACTGCTGAGTATTCCCCTTTCAAGGGAGGTTACACCCAAAGGGCTAGAAGGCGAGA GITTAGGITGITTIGGIAGCGITGIAGTAGGGAGITTCGGGAIAITTIGAATTGICGITTA CCCTGTCTCTACAAGTCCGGAGCACTGGACGAGGCAGCTGCGTACCAGAGTCGCGACT CGGTGTCCATGGGCTGGGTGTGGAGGCGTTGGAGCATCTGAGTCCAGGGGAACAGCTTC 2120 CCAGGCAGCAGCAGCAGCAGCAGGGTGATGGTTCTCCCCCAAGCCCATCGTAGAGGCC CCACAGGCTACCTGGTCCTGGATGAGGAACAGCAACCTTCACAGCCGCAGTCGGCCCTGG 2240 AGTGCCACCCGAGAGGTTGCGTCCCAGAGCCTGGAGCCGCCGTGGCCGCCAGCAAGG TGTCCCTGCTGGGCCCCCACTTTCCCCGGCTTAAGCAGCTGCTCCGCTGACGTTAAAGACA TCCTGAGCGAGGCCAGCACCATGCAACTCCTTCAGCAACAGCAGCAGGAAGCAGTATCCG ATTACTTAGGGGGCCACTTCGACCATTTCTGACAACGCCAAGGAGTTGTGT.AGGCAGTGT ATTATTTAGGGGGTATTTCGATTATTTTTGATAACGTTAAGGAGTTGTGTAAGGTAGTGT GGGGGGATTGCATGTACGCCCCACTTTTGGGAGTTCCACCGGGTGTGCGTCCACTTCTT GGCTGGGAAGGGTCTACCCTCGGCCGCGTCCAAGACCTACCGAGGGCTTTCCAGAATC GCGCAGCACCTCCCGGCGCCCAGTTTGCTGCTGCTGCAGCAGCAGCAGCAGCAGCAGCAGCAGC TTAGGTAGTAGTAGTAGTAGGGTGGGTGGTTTTTTTTTAGGTTTTTATCGTAGAGGTT **AAGGCAGCAGCAGGGAGAGCGAGGGAGGCCTCGGGGGCTCCCACTTCCTCCAAGGACA** GCGTAGTATTTTCGGCGTTAGTTTGTTGTTGTT------GT TGTTCCAGAGCGTGCGCGAAGTGATCCAGAACCCGGGCCCCAGGCACCCAGAGGCCGGA 2840 2780 1484 2300 2360 1004 2420 1064 2480 1124 2540 1184 2600 2660 2060 2180 1880 1940 2000 764 884 944 548 809 899 704 g qq g ŏ g δ a ŏλ g ò qq g δ ğ δ g QY g ŏ Db Qγ qq Qy g ò g ŏ ò g g ŏ g

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                                                                                                                                                                                                                                                                                                                                     The present invention relates to a method for analysis of inactivation of x chromosome. The method comprises analysing methylation of human androgen receptor (HUMARA) gene (the present sequence) by a methyl-specific polymerase chain reaction (PCR). The PCR primers amplify the base sequence of the region containing repeated number of polymorphism of CAG of the HUMARA gene, specific to methylation of the cytosine base at the 199th, the 203rd and the 206th or the 296th position of the present sequence. The method is useful for the detection of uniformity of cell growth, that is clonality.
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                                                                                                androgen
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98.7%; Pred. No. 1.8e-284;
ive 0; Mismatches 4;
                                                                                                 inactivation;
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                                                                                                 Human: PCR primer; X chromosome
methylation; HUMARA; ds.
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                     DNA; 1810
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Matches 1767; Conservative
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                    AAF84342 standard;
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                                                                                                                                                                                                                                                                                                                2481 AGGCAGCAGCAGCGGGAGGCGAGGGCCTCGGGGGCTCCCACTTCCTCCAAGGACAA
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3261 CGGCGGCGGCGGCGGCGGCGGCGGCGGCGGGGGGGGGTGTAGCCCCTACGGCTA 3320
                                                3321 CACTCGGCCCCTCAGGGGCTGGCGGGCCAGGAAAGCGACTTCACCGCACCTGATGTGTG 3380
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